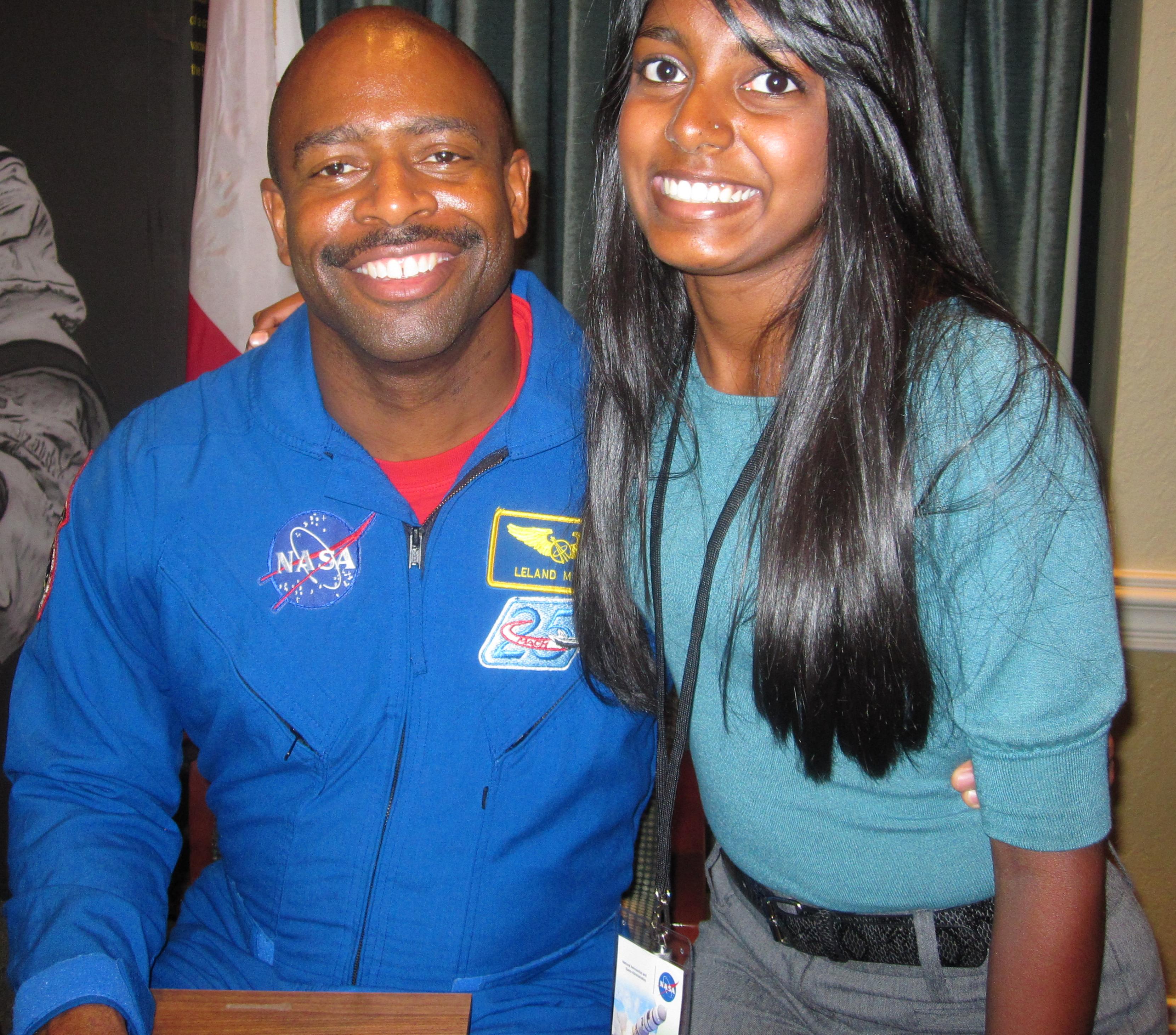


Abstract

MuRGE (Mutualism in a Reduced Gravity Environment) is a NASA flight-research experiment to investigate the microgravity effects associated with cell-cell communication and beneficial microbe-host interactions using a plant-fungal model system. This investigation will use a clinostat, an instrument that slowly rotates the plants to negate the effects of gravitational pull on plant growth (gravitropism) and development, to simulate microgravity. I will be using the endophytic fungus *Piriformospora indica* (Pi) and the model plant species *Arabidopsis thaliana* (At). *P. indica* has been shown to colonize roots of various plant species, including *A. thaliana*, and to increase plant growth and resistance to stress. The fungus has the ability to grow from spores or in axenic cultures without the presence of a host. *P. indica* spores and *P. indica* extract will be used to inoculate Arabidopsis seeds germinated on a clinostat in order to determine if simulated microgravity affects the interaction between the fungus and its plant host.









NASA INSPIRE

Karishma Patel
Pre-college Intern
Kennedy Space Center, FL
Summer 2010

About Me

- ▶ Karishma Patel
- ▶ Hometown: Calhoun, Georgia
- ▶ Georgia Institute of Technology, Class of 2014
- ▶ Hobbies: Running, playing the trumpet, hiking



Overview

- ▶ Work Overview
- ▶ MuRGE
- ▶ Exposure to
- ▶ Work Related Activities
- ▶ Non-Work Related Activities
- ▶ Lessons Learned
- ▶ Lasting Impact
- ▶ Future Plans
- ▶ Acknowledgements

Work Overview

- ▶ Mentor: Michael Roberts
- ▶ Space Life Sciences Lab (SLL)
- ▶ Dynamac Corporation
- ▶ NASA Exploration Life Support Program
- ▶ NASA-KSC Engineering Directorate (NE-S)

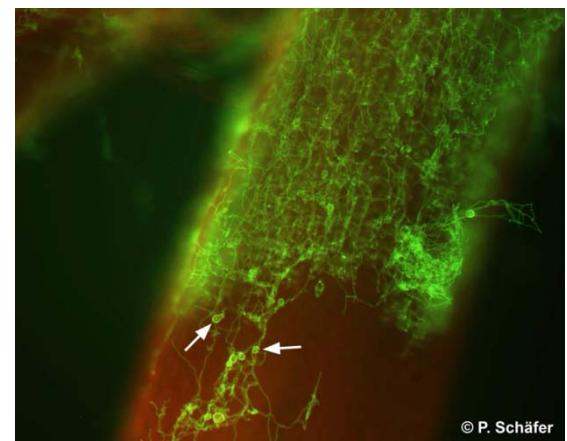


MuRGE (Mutualism in a Reduced Gravity Environment)

- ▶ Mutualism is the way two organisms biologically interact where each individual derives a fitness benefit.
- ▶ Microgravity affects the physiology of organisms and the way they interact with each other and their environment. This may result in increased risk of infections.
- ▶ MuRGE is a flight experiment to study the effect of microgravity on the interaction between a plant (*Arabidopsis thaliana*) and a fungus (*Piriformospora indica*).



Arabidopsis thaliana (thale cress)



Piriformospora indica
(green hyphae attached to root)
© P. Schäfer

MuRGE

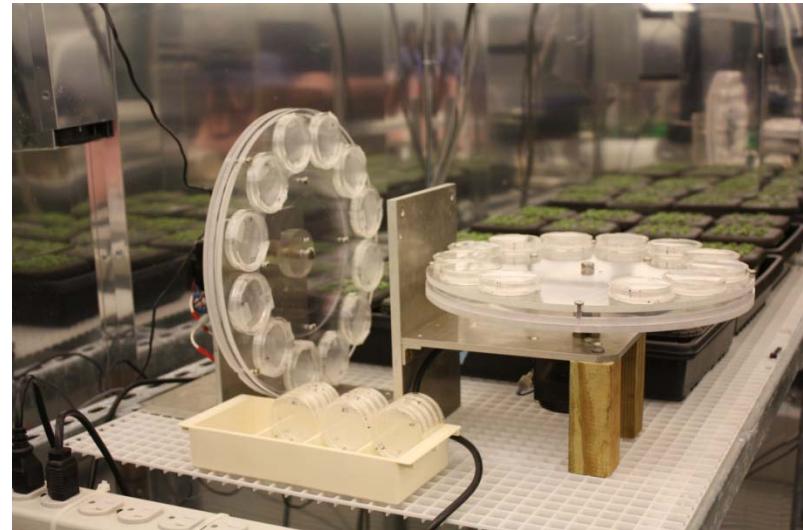
► Why study plants and fungi in space?

- Microorganisms (bacteria, archaea, fungi, viruses) are part of the spacecraft human environmental control and life support system. The microbes go where we go.
- Plants can be a part of the life support system as well to provide fresh food, improve air quality (removing CO₂ and VOCs while providing O₂), and recycle wastewater.
- Living organisms (whether human, plant or microbe) are affected by living in reduced gravity. In some cases, this can result in an increased risk of infection and disease.
- In normal gravity, the fungus *P. indica* increases the rate of plant growth and resistance to environmental stress.

MuRGE

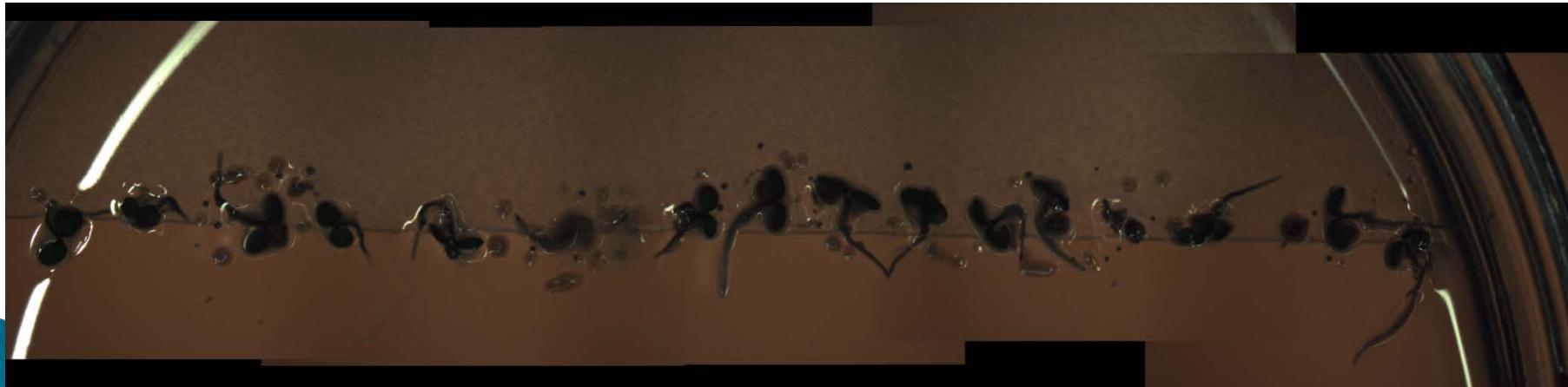
- ▶ Clinostats
 - Simulate the effects of reduced gravity on organisms by rotating them at low speed

- ▶ Controlled Environment Chamber (CEC)
 - 18:6hr Light:Dark, 400ppm CO₂, 20°C, and 50% RH (relative humidity)



MuRGE

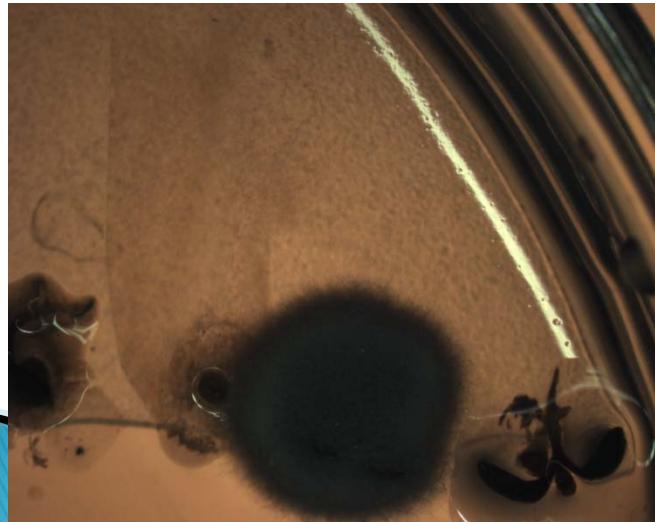
- ▶ 36 agar plates (18 seeds/plate)
 - 12 horizontal, 12 vertical, 12 control
 - 3 At only
 - 3 Pi only
 - 3 At+Pi spores
 - 3 At+Pi extract



At and Pi extract (7x)- Day 4

MuRGE

- ▶ Null hypothesis: *P. indica* will increase the growth of *A. thaliana* in both normal gravity and simulated microgravity.
- ▶ 2 experiments
 - Contaminated
 - No Pi growth
- ▶ Pi did not increase plant growth in normal gravity or simulated gravity



Exposure to...

Biolog

- ▶ Microbial species identification
 - 96-well plate
 - Yeasts, fungi, bacteria
 - Transmittance
 - Incubator- 30/37°C



Other

- ▶ DNA extractions
- ▶ Acridine Orange (AO) Stains
- ▶ Microscopy
- ▶ Spectroscopy



Work Related Activities

- ▶ Seminars
- ▶ VAB
- ▶ Intern socials



Non-Work Related Activities



- ▶ Harry Potter Land
- ▶ Beach
- ▶ SAK comedy club



Lessons Learned

- ▶ Be courteous and on time
- ▶ Be flexible
- ▶ Be prepared for anything
- ▶ Don't be afraid to ask questions

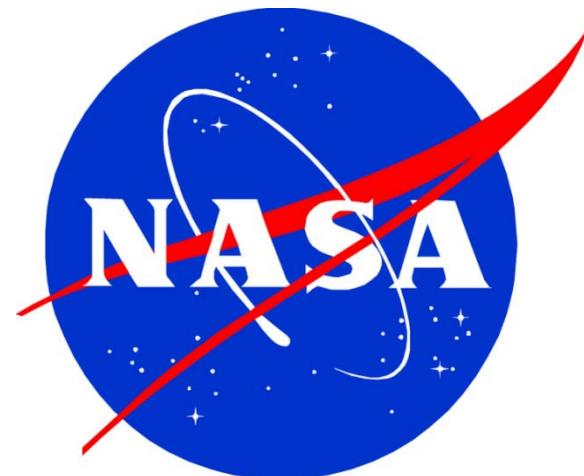
Lasting Impact

- ▶ Lab skills
- ▶ Work experience
- ▶ Appreciation for NASA
- ▶ Understanding of workplace ethics
- ▶ Preparation for college



Future Plans

- ▶ Georgia Institute of Technology
- ▶ Major: Biology
- ▶ Future job at NASA



Acknowledgements

- ▶ Michael Roberts
- ▶ Michele Birmele
- ▶ Priscilla Moore
- ▶ Amber Wade
- ▶ Jim Gerard
- ▶ Luke Roberson
- ▶ NASA INSPIRE

INSPIRE Pre-College Internship
Kennedy Space Center
Mutualism in a Reduced Gravity Environment (MuRGE)
Patel, Karishma
July 29, 2010

Reviewed by:

Michael S. Roberts, Ph.D.

Mail Code DYN-3

Dynamac Corporation

Engineering Directorate, Surface Systems Office (NE-S)

Abstract

MuRGE (Mutualism in a Reduced Gravity Environment) is a NASA flight-research experiment to investigate the microgravity effects associated with cell-cell communication and beneficial microbe-host interactions using a plant-fungal model system. This investigation will use a clinostat, an instrument that slowly rotates the plants to negate the effects of gravitational pull on plant growth (gravitropism) and development, to simulate microgravity. I will be using the endophytic fungus *Piriformospora indica* (Pi) and the model plant species *Arabidopsis thaliana* (At). *P. indica* has been shown to colonize roots of various plant species, including *A. thaliana*, and to increase plant growth and resistance to stress. The fungus has the ability to grow from spores or in axenic cultures without the presence of a host. *P. indica* spores and *P. indica* extract will be used to inoculate Arabidopsis seeds germinated on a clinostat in order to determine if simulated microgravity affects the interaction between the fungus and its plant host.

Introduction

Mutualism can be described as interactions between different organisms that are beneficial for both partners. One common effect of mutualism is increased growth of one or both organisms by increasing access to nutrients. Microgravity affects the physiology of organisms and the way they interact with one another and their environment. Plants and fungi go wherever humans go and are therefore a part of spacecraft environmental control and life support systems. It is necessary to study the effects of microgravity on microorganisms so we can be prepared for long duration missions in space. MuRGE will test the interaction between At and Pi in both normal gravity and microgravity environments. The null hypothesis is that *P. indica* will increase the growth of *A. thaliana* in both normal gravity and simulated microgravity.

Materials and Procedures

This experiment will germinate seeds from *A. thaliana* (L.) Heynh wild-type ecotype Columbia on a solid agar media consisting of 2.2 g of Murashige and Skoog salts (Murashige and Skoog, 1962), 0.5 g of MES buffer, 2.5 g of sucrose, and 1 mL of 1,000x Gamborg vitamins (Sigma-Aldrich, St. Louis) per liter at pH of 5.75. Phytagel (Sigma) is added to a concentration of 0.8% (w/v) and autoclaved. After autoclaving, 5 mL of the media is aliquoted into a sterile 60 mm Petri plate (Millipore Microbiological Dishes, 47mm). Each Petri plate will be planted

with 18 sterilized *Arabidopsis* seeds per plate oriented in a single row across the plate on the solid surface. For *P. indica* spore treatments, *A. thaliana* seeds are coated with *P. indica* spores immediately prior to planting by immersing the seeds for 2 minutes in ~50 µL of a spore suspension containing $\sim 1 \times 10^4$ spores mL⁻¹ dI H₂O. For *P. indica* extract treatments, *A. thaliana* seeds are coated with *P. indica* extract immediately prior to planting by immersing the seeds for 2 minutes in ~50 µL of the extract.

Four slow-rotation clinostats rotating at 1 rpm, each holding 12 60-mm Petri plates arranged in 4 sets of 3, will be used during the experiment. Three Petri plates will contain only *Arabidopsis thaliana* (Treatment A), three plates will contain only *P. indica* (Treatment B), three will combine *A. thaliana* and *P. indica* spores (Treatment C), and a third set of plates will contain *A. thaliana* and an extract of *P. indica* (Treatment D). A third set of 12 60-mm Petri plates will be placed on the shelf as a no-centrifugal force control. Each Petri plate will be prepared in duplicate and one plate corresponding to each treatment will be covered in aluminum foil as an etiolated (dark) treatment. Two clinostats will be oriented vertically to rotate the Petri plates perpendicular to the Earth's surface (Simulated Microgravity) and the other clinostats will be oriented horizontally to rotate the Petri plates parallel to the Earth's surface (Normal Gravity Control). Petri plates loaded into the clinostats and the static controls will be placed in a Controlled Environment Chamber (CEC 15) under the following growth conditions: 18:6 Light:Dark, 300 µmol m⁻² sec⁻¹, 400 ppm CO₂, 20°C, and 50% RH (relative humidity).

The growth of the plants will be monitored by measuring the length of roots and shoots of the *Arabidopsis* plants on the Petri plates and measuring the fresh-weight biomass. A single Petri plate will be harvested from each treatment at each of three four-day time intervals spanning twelve days. At each three-day interval, one plate from each treatment will be removed from the clinostat and the plants harvested for measurement under an Olympus SZX-12 zoom microscope.

TREATMENTS (18 seeds per Petri plate; 3 Petri plates per treatment)

A. <i>A. thaliana</i>	CTL ; HC ; VC	3 plates/treatment = 9 plates
B. <i>P. indica</i>	CTL ; HC ; VC	3 plates/treatment = 9 plates
C. <i>A. thaliana</i> and <i>P. indica</i> spores	CTL ; HC ; VC	3 plates/treatment = 9 plates
D. <i>A. thaliana</i> and <i>P. indica</i> extract	CTL ; HC ; VC	3 plates/treatment = 9 plates 36 plates incubated in light 36 plates incubated in dark

	DOE4	DOE8	DOE12
Control	CTL: A1/B1/C1/D1	CTL: A2/B2/C2/D2	CTL: A3/B3/C3/D3
Horizontal Clinostat	HC: A1/B1/C1/D1	HC: A2/B2/C2/D2	HC: A3/B3/C3/D3
Vertical Clinostat	VC: A1/B1/C1/D1	VC: A2/B2/C2/D2	VC: A3/B3/C3/D3

Results

1st Experiment-Day 4

Plate	# of seeds	Germinated	Orientation					Notes
			Up	Down	Left	Right	Unknown	
A1 HC	18	14	6	4	2	2		
A1 N	18	17	15	2				
A1 VC	18	13	2	10	1			
B1 HC	18							16 spots of Pi (fungal contaminant?)
B1 N	18							Cannot distinguish individual spots, fungal growth along center
B1 VC	18							Cannot distinguish individual spots, fungal growth along entire line, one large growth on right
C1 HC	18	17	7	2	4	4		Spores and fungal growth not visible
C1 N	18	15	10	3	2			14 seem to have spores, 2 of those overtaken by Pi
C1 VC	18	16	5	5	3	3		11 visibly affected by Pi, others not visible
D1 HC	18	15	5	3	4	3	3	Plate had bubbles, unable to determine if affected by Pi
D1 N	18	13	11			2		Longest roots, all affected by Pi, 1 overtaken by Pi
D1 VC	18	16	3	10	1	2		Fungal growth visible across entire line, 1 overtaken

The data above was collected from the first set of plates, which contained eighteen seeds each. In most cases, more than three-fourths of the seeds germinated. In the neutral plates, a majority of the plants sprouted upwards. However, a few of the seeds that had recently germinated grew in other directions. The plates that were placed on the horizontal clinostat had similar results to one another. A majority of plants in each plate grew upwards. The plates on the vertical clinostat had a majority of seeds that grew downwards. The Pi growth was undistinguishable from the fungal contamination.

1st Experiment-Day 8

Plate	# of seeds	Germinated	Orientation					Notes
			Up	Down	Left	Right	Unknown	
A2 HC	18	17	14	3				
A2 N	18	17	17					
A2 VC	18	18	4	9	8	2		
B2 HC								10 visible colonies of contaminant (?)
B2 N								2 types of fungal contaminant
B2 VC								2 types of fungal contaminant
C2 HC	18	17	5	2	4		6	Pi/ contaminant has overtaken At-unable to observe roots/orientation
C2 N	18	16	6	5			5	Pi/ contaminant has overtaken At-unable to observe roots/orientation
C2 VC	16?	14	4	5	2	3		Plate contaminated, 5 overtaken
D2 HC	18	17						Plate dropped? All plants are loose and floating
D2 N	18	17	17					Bubbles around left side of plate, contaminate on right side
D2 VC	18	18	8	5	1		4	Many are shriveled, germinated at dif. times, newly germinated have been overtaken by contaminant

The data above was collected from the second set of plates from the first experiment. All of the plates, except one, showed to have eighteen seeds. A majority of those seeds germinated. In the *Arabidopsis* only and *Arabidopsis + P. indica* extract, all of the germinated seeds grew upwards.

However, the *Arabidopsis* + *P. indica* spores had a randomly distributed amount of plants in all directions. The plates that were placed on the horizontal clinostat had a majority of plants that grew upwards. One plate had floating seeds, which could have resulted from the plate being dropped. The plates on the vertical clinostat had a randomly distributed amount of seeds growing in all directions. The direction and rate of Pi growth was unable to be quantified because of the presence of fungal contaminant(s).

2nd Experiment-Day 4

Plate	# of seeds	Germinated	Orientation				Notes
			Up	Down	Left	Right	
A1 HC	18	18	9	4	4	1	
A1 N	18	15	15				
A1 VC	18	15	5	10			
B1 HC							1 distinguishable spot
B1 N							Pi not visible
B1 VC							Pi not visible
C1 HC	18	15	7	6		2	Pi not visible
C1 N	18	16	16				Root hairs visible
C1 VC	18	15	5	8	2		
D1 HC	18	14	7	4			3 A couple of roots are interwoven-unable to determine orientation
D1 N	18	18	18				Longest roots, root hairs somewhat visible
D1 VC	18	15	6	6		3	Pi not visible

The data above was collected from the first set of plates from the second experiment. In most cases, more than three-fourths of the seeds germinated. In the neutral plates, all of the seeds that germinated grew upwards. The plates that were placed on the horizontal clinostat had a majority of the plants grew upwards. The plates on the vertical clinostat had a majority of seeds that grew downwards. The Pi did not show any visible growth on these plates.

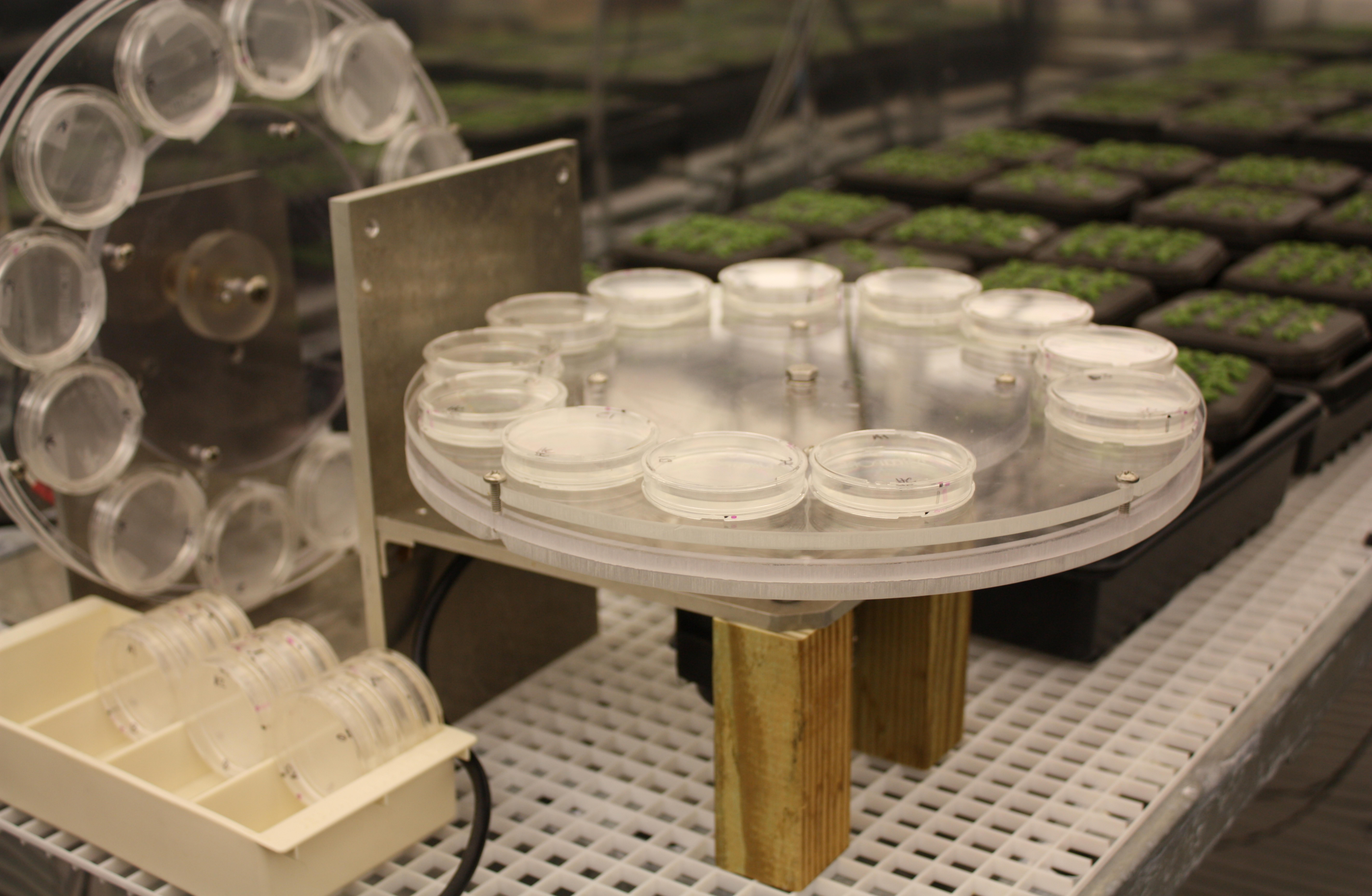
Discussion

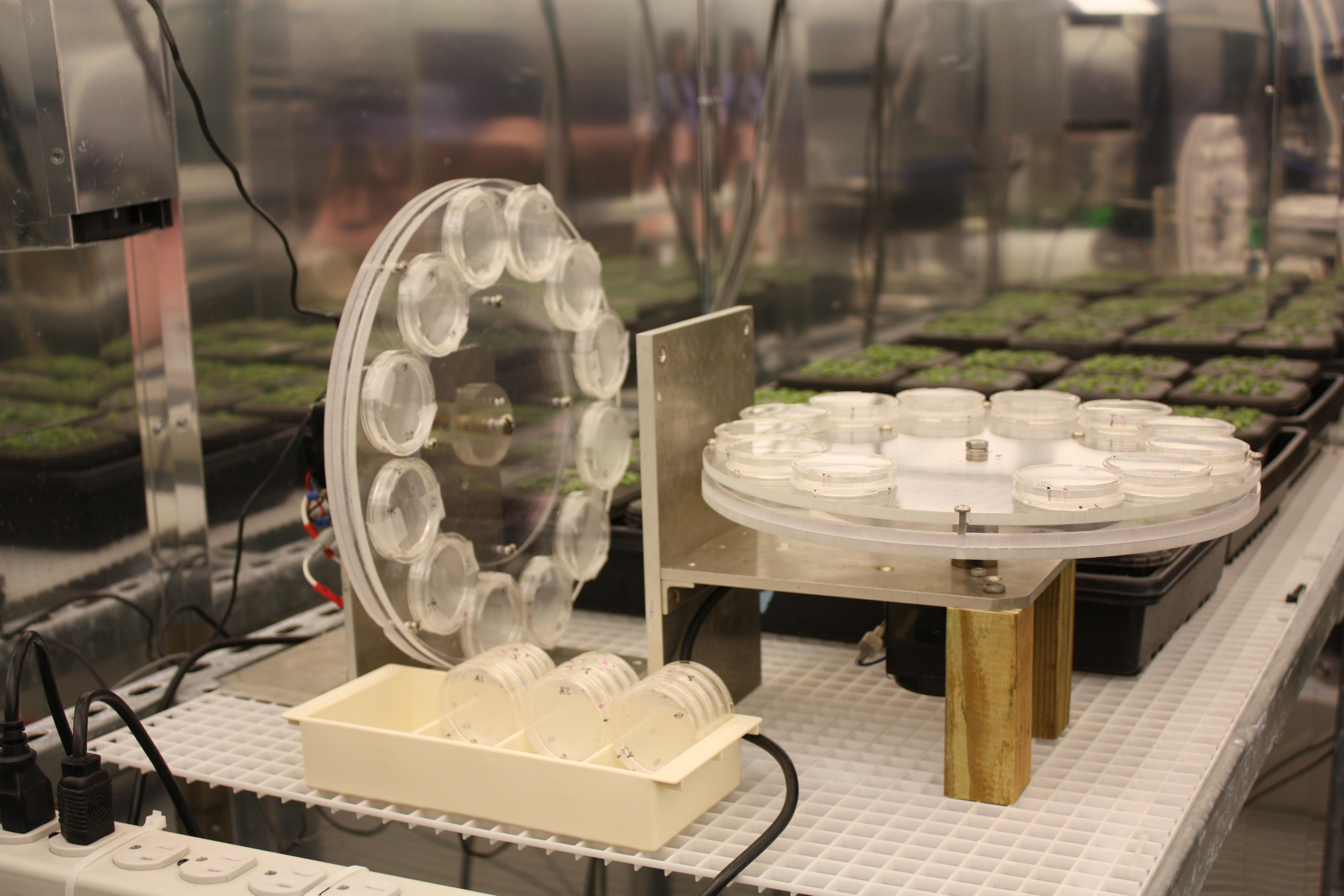
MuRGE investigated the microgravity effects associated with cell-cell communication and beneficial microbe-host interactions using a plant-fungal model system. This investigation used a clinostat to simulate the effects of microgravity. The plant *Aribidopsis thaliana* and the fungus *Piriformaspora indica* were used to investigate these microbe-host interactions.

Two experiments were completed using the clinostats. While collecting data from the day four plates, a contaminant was spotted on the plates. The spores were the likely source of this. The inconclusive experiment resulted in starting a second experiment. Unfortunately, there was no visible spore growth in the first eight days of the experiment. Due to time restraints of my internship, I was unable to collect data for the day twelve plates of my second experiment. As a result of two inconclusive experiments, I was unable to determine the affects of Pi on At in both normal gravity and microgravity.

Acknowledgements

I would like to thank my mentor, Michael Roberts, for all the time and support he gave me this summer and Michele Birmele for teaching me lab techniques and procedures. I would also like to thank all of the other staff at the Space Life Sciences Lab for making my summer experience an amazing one.





CEC 15

Environmental
Growth Chambers

CAUTION

CALIBRATE
BEFORE USE
NASA

13.0
50
14.0E

TC2 TEMP & RH BOARD
ID (PLOC 20)=12

12.50
11.6
14.0E

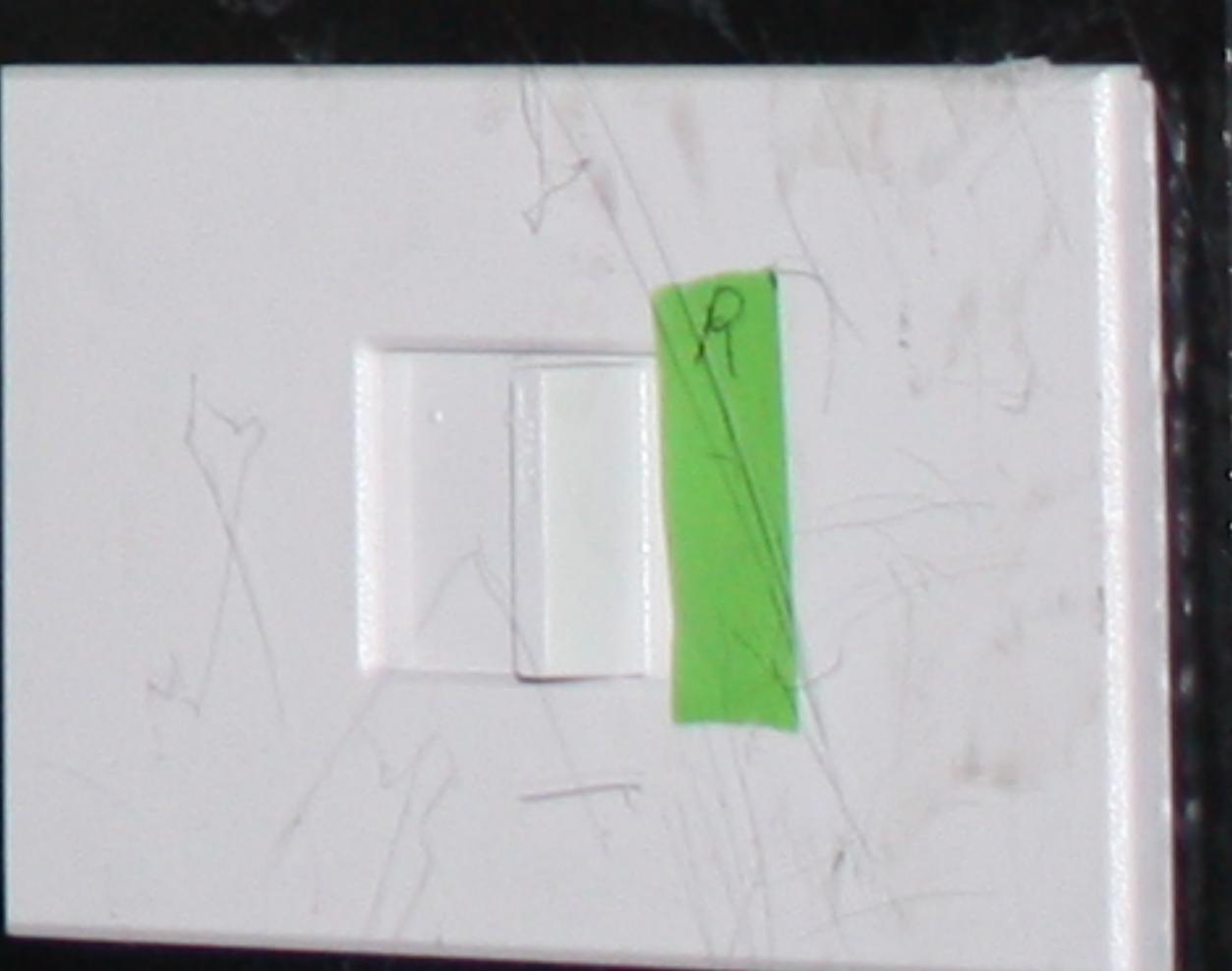
TC2 CO2 & LIGHT BOARD
ID (PLOC 20)=13

SSR 1=LIGHT CAP 1 ENABLE
SSR 2=LIGHT CAP 2 ENABLE

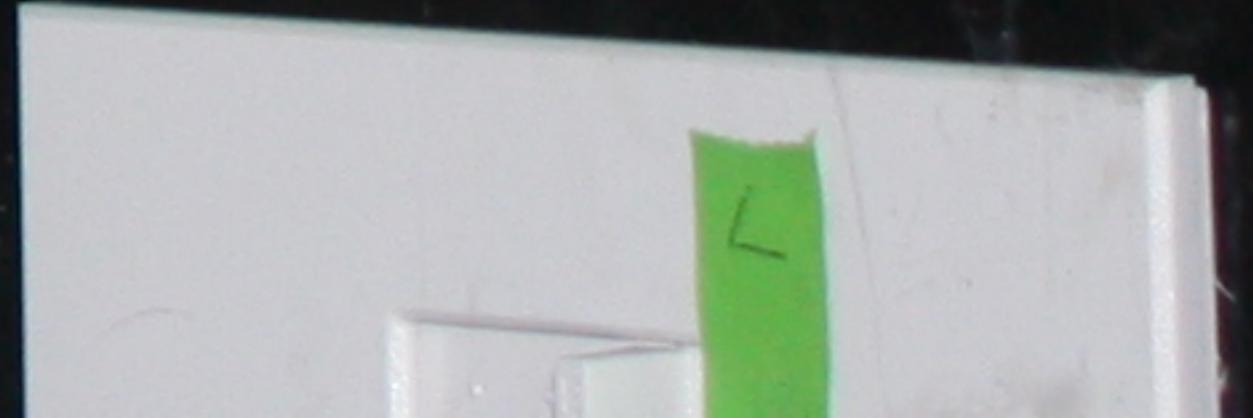
▲
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EGC
TC2
ENTER

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EGC
TC2
ENTER

LIGHTCAP #1

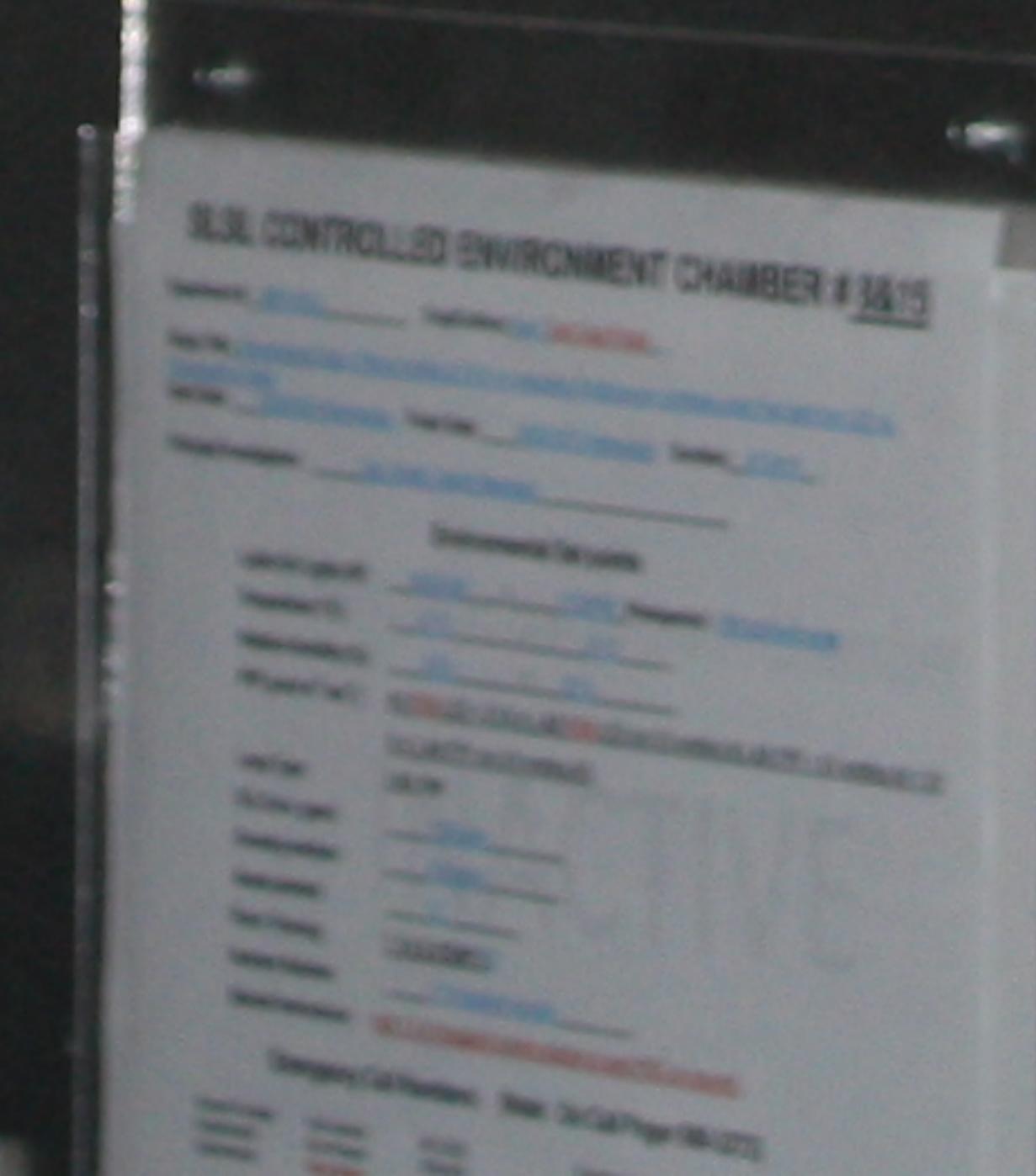


LIGHTCAP #2



Environmental
Growth Chambers
PO Box 390, Chagrin Falls, Ohio 44022
USA

CEC 15



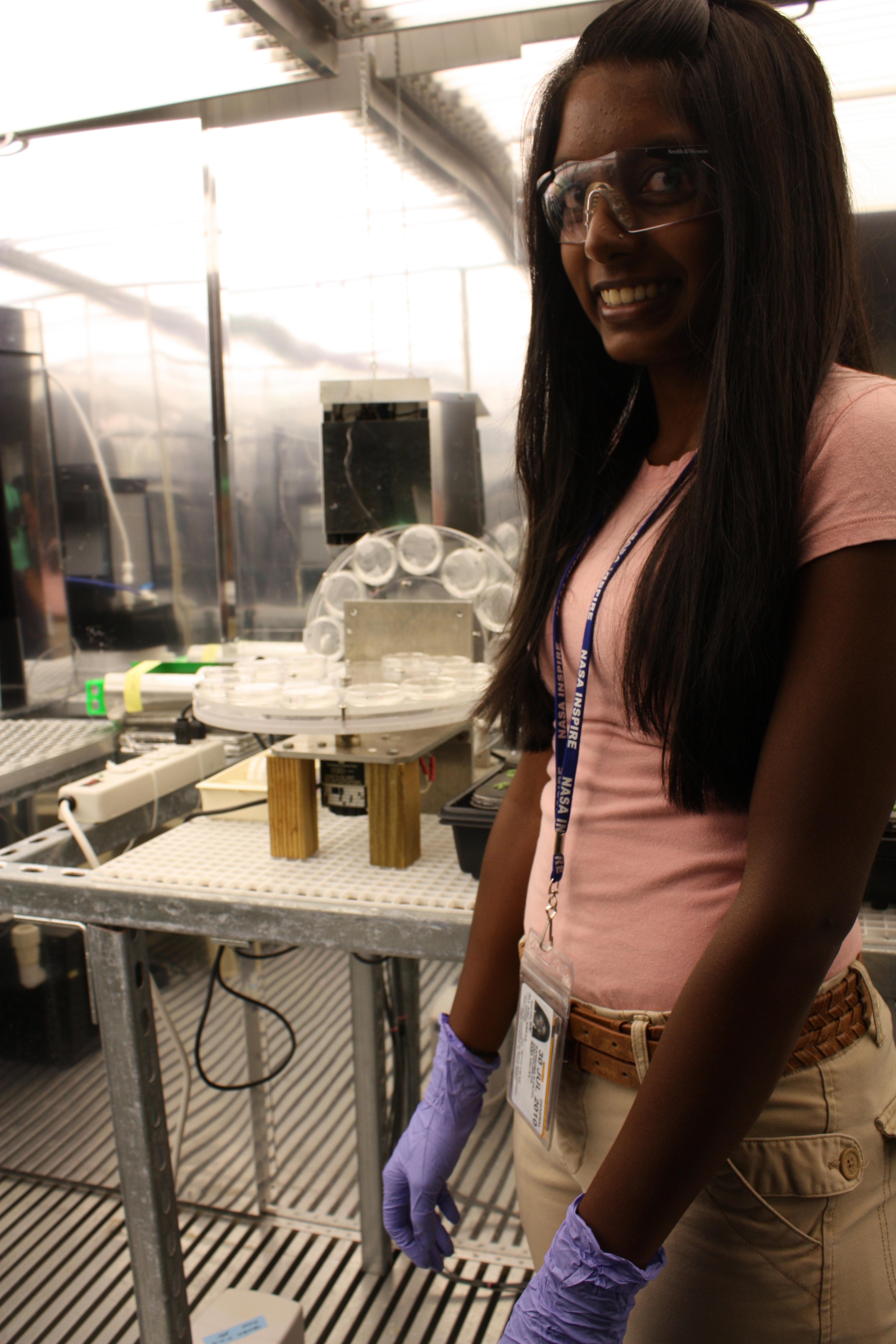
The image shows two identical control panels for TC2 modules, one above the other. Each panel has a digital display showing red numbers. The top panel displays 13.4, 50, and 140E. The bottom panel displays 1245, 116, and 140E. Both panels feature a 'CALIBRATE BEFORE USE' label with NASA certification at the top. They include a vertical stack of buttons: an 'OUT' button, a small rectangular button labeled 'SSR', and four checkboxes labeled RAMP, LOCK, DEFROST, and ALARM. To the right of these are four more buttons: RUN, EDIT, MOD, and ENTER, each with a corresponding up and down arrow icon. A label 'TC2 TEMP & RH BOARD ID (PLOC 20)=12' is located below the top panel's display. A label 'TC2 CO2 & LIGHT BOARD ID (PLOC 20)=13' is located below the bottom panel's display. At the very bottom, a box contains the text 'SSR 1=LIGHT CAP 1 ENABLE' and 'SSR 2=LIGHT CAP 2 ENABLE'.

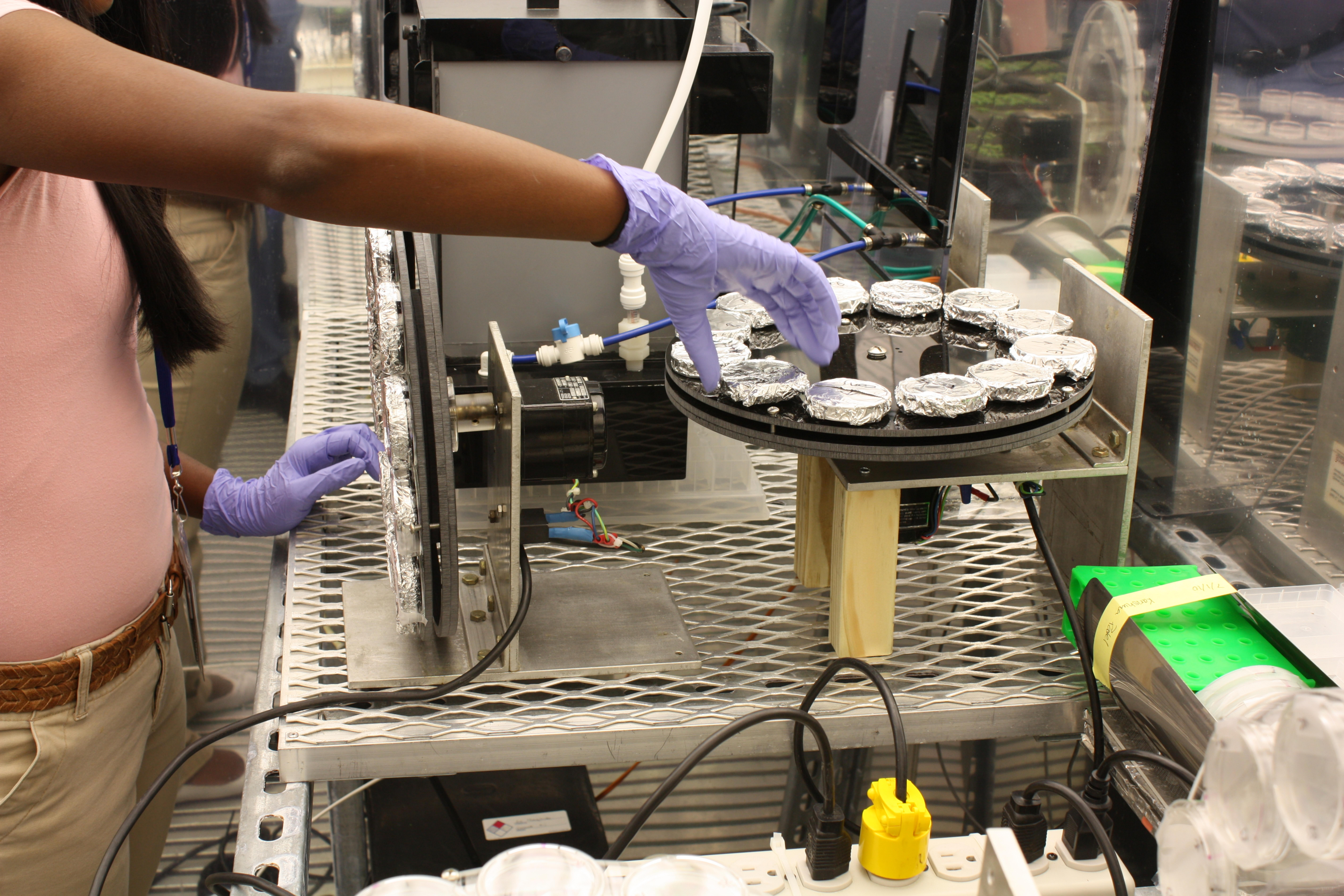


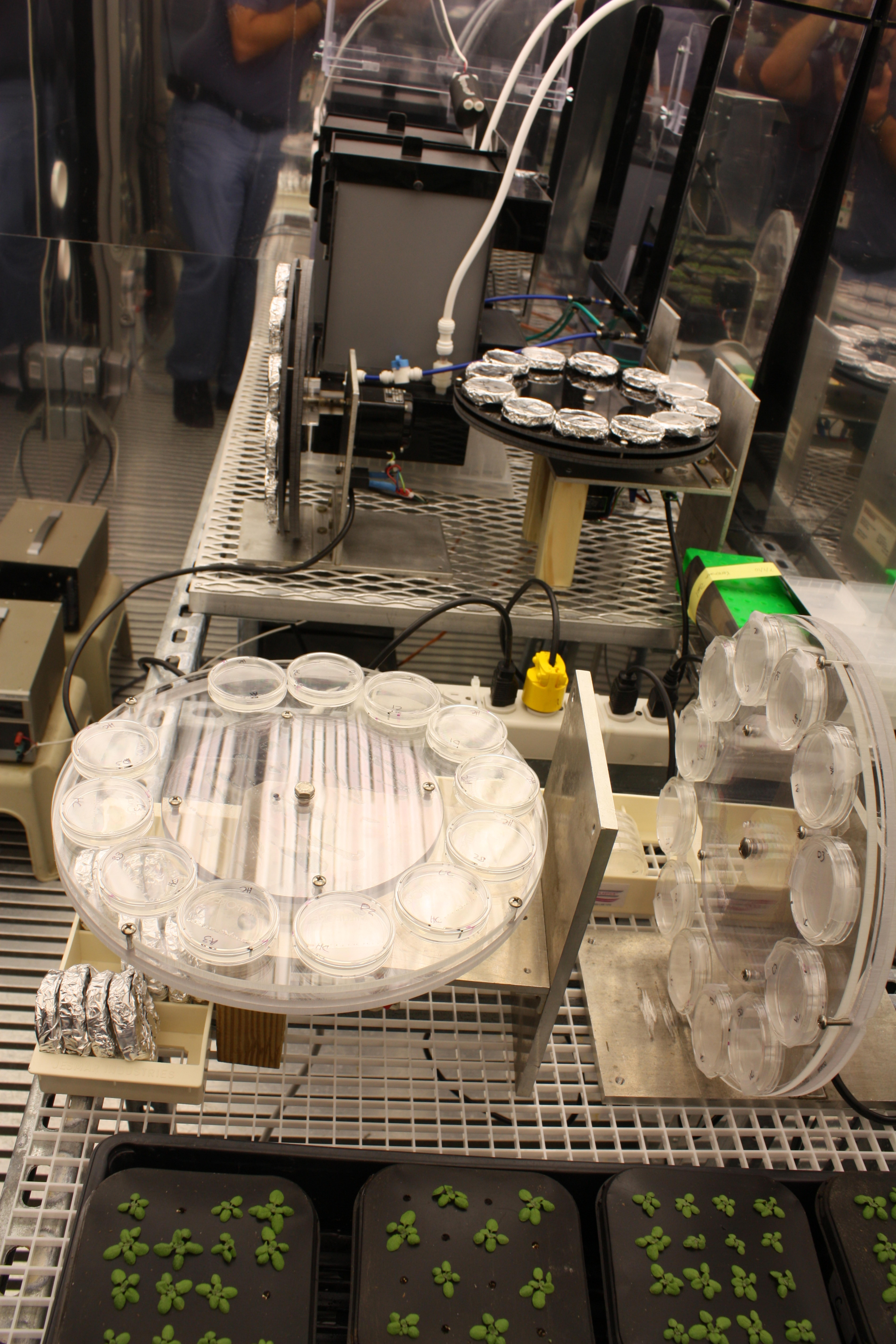


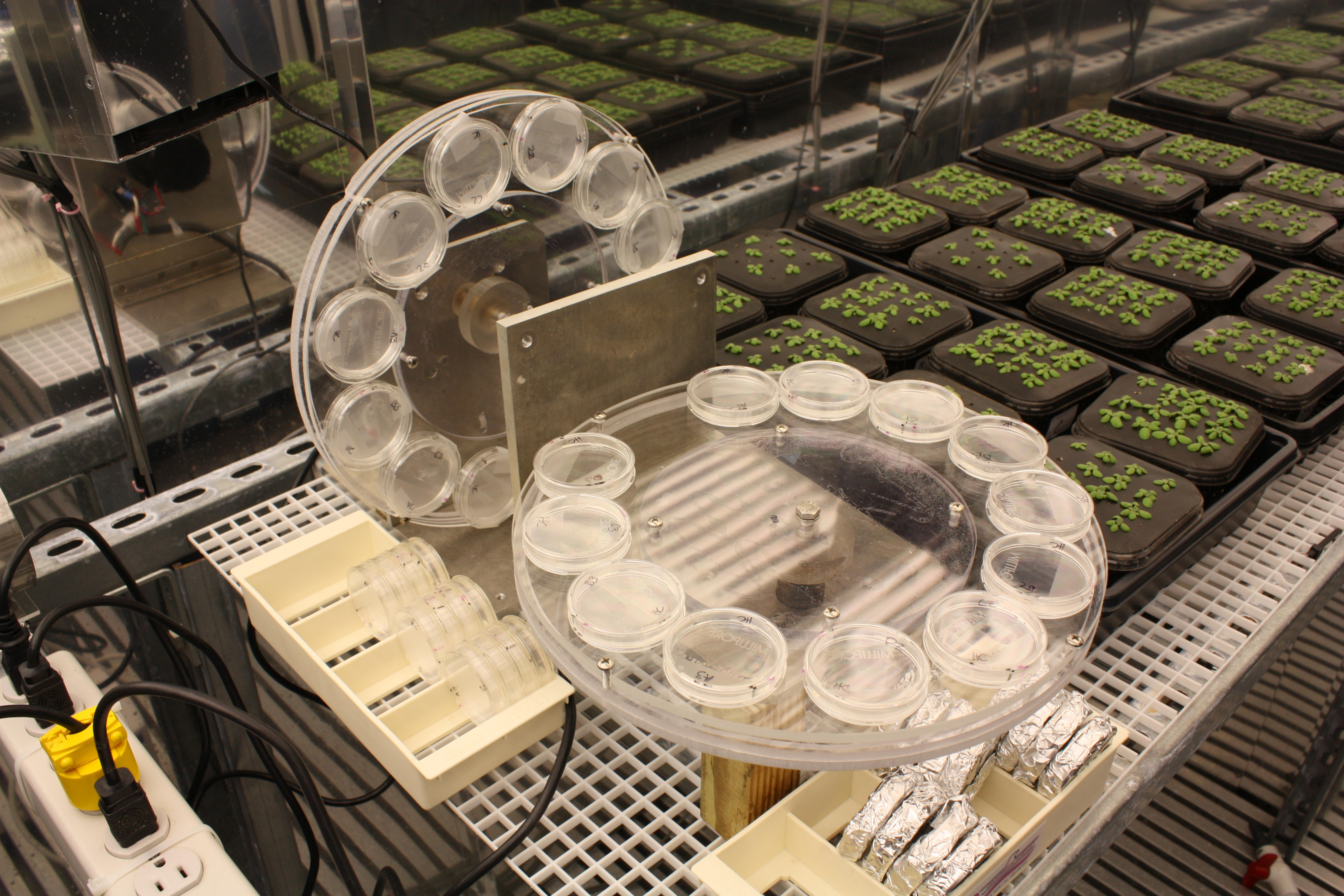






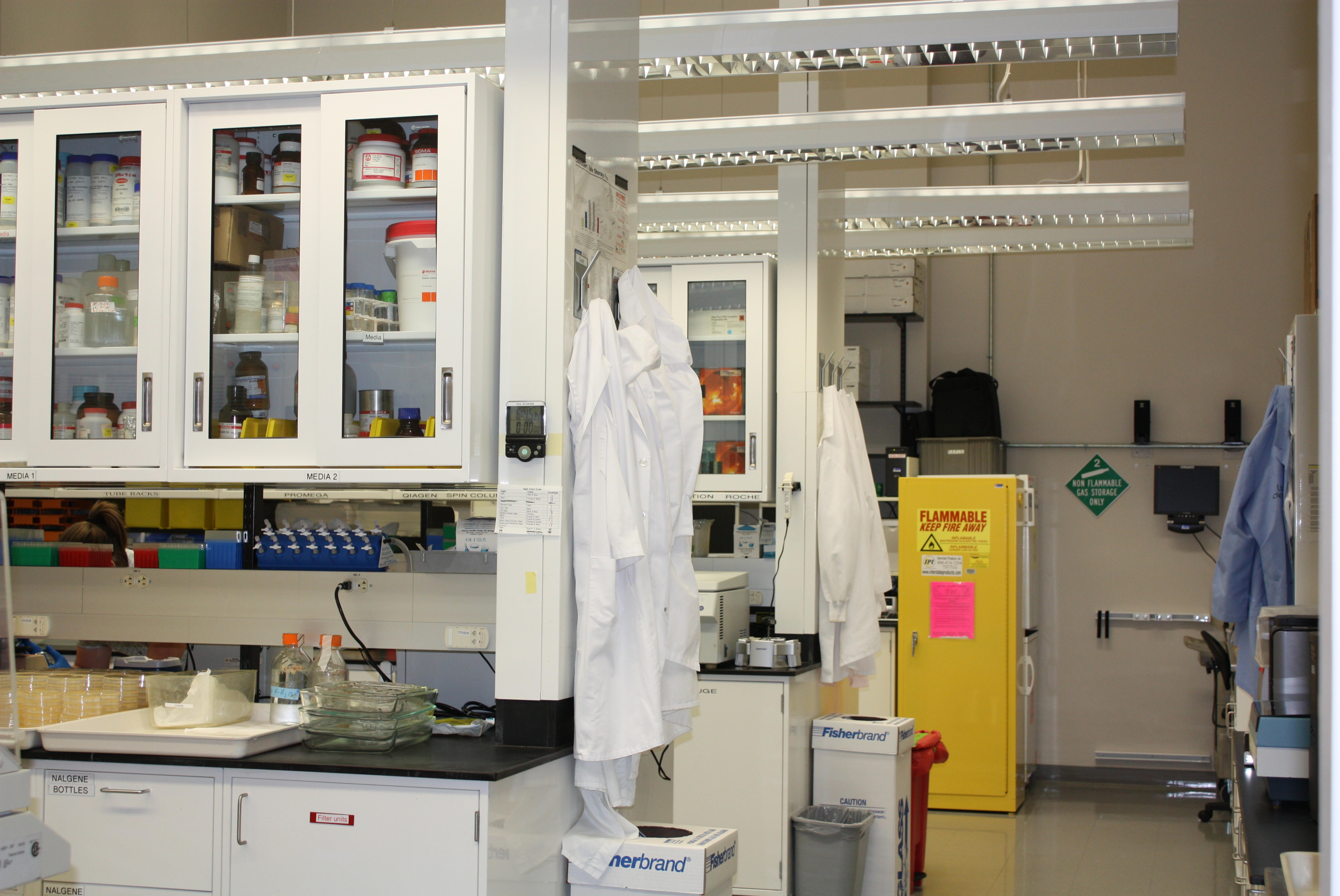






















15

16

